



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification⁶ : A61K 9/70</p>	<p>A1</p>	<p>(11) International Publication Number: WO 96/19205 (43) International Publication Date: 27 June 1996 (27.06.96)</p>
<p>(21) International Application Number: PCT/US95/16498 (22) International Filing Date: 8 December 1995 (08.12.95) (30) Priority Data: 08/361,800 21 December 1994 (21.12.94) US (71) Applicant: THERATECH, INC. [US/US]; Suite 100, 417 Wakara Way, Salt Lake City, UT 84108 (US). (72) Inventors: EBERT, Charles, D.; 1515 South Canterbury Drive, Salt Lake City, UT 84108 (US). HEIBER, Werner; 1569 So. 2300 East, Salt Lake City, UT 84108 (US). GOOD, William, R.; 7730 So. Quicksilver, Salt Lake City, UT 84121 (US). VANKATESHWARAN, Srinivasan; 2411 Emerson Avenue, Salt Lake City, UT 84108 (US). (74) Agents: KONSKI, Antoinette, F. et al.; Morrison & Foerster, 755 Page Mill Road, Palo Alto, CA 94304-1018 (US).</p>		<p>(81) Designated States: AU, CA, CN, JP, KR, MX, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
<p>(54) Title: TRANSDERMAL DELIVERY SYSTEM WITH ADHESIVE OVERLAY AND PEEL SEAL DISC</p> <div data-bbox="354 1197 1156 1570"> </div> <p>(57) Abstract</p> <p>A device (10) for administering an active agent to the skin or mucosa of an individual comprising a laminated composite of an adhesive overlay (26), a backing layer (14) underlying the central portion of the adhesive overlay, an active agent-permeable membrane (16), the backing layer and membrane defining a reservoir (12) that contains a formulation of the active agent, a peel seal disc (20) underlying the active agent-permeable membrane, a heat seal (22) about the periphery of the peel seal disc, the active agent-permeable membrane and the backing layer and a removable release liner (24) underlying the exposed overlay and peel seal disc. The adhesive layer is above and peripheral to the path of the active agent to the skin or mucosa and is protected from degradation by the components of the reservoir by a multiplicity of heat seals. The peel seal disc protects against release of the active agent-containing reservoir and the release liner protects the adhesive from exposure to the environment prior to use.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LJ	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

5 TRANSDERMAL DELIVERY SYSTEM WITH ADHESIVE OVERLAY AND
 PEEL SEAL DISC

Technical Field

10 This invention is in the field of transdermal
 and transmucosal administration of active agents (drugs).
 More particularly, it relates to a device for achieving
 such administration that has an active agent-containing
 reservoir and an adhesive layer for affixing the device
15 to the skin or mucosa. The adhesive layer is above and
 peripheral to the path of the active agent to the skin or
 mucosa and is protected from degradation by the
 components of the reservoir by a multiplicity of heat
 seals. A peel seal disc protects against release of the
20 active agent-containing reservoir and a release liner
 protects the adhesive from exposure to the environment
 prior to use.

Background of the Invention

25 Devices for administering drugs through the
 skin or mucosa have been described in many patents.
 These devices are usually of two types: matrix systems
 and liquid reservoir systems. Both systems are laminated
 composites that have, from basal to top, a peelable
30 release liner, a pressure sensitive adhesive layer for
 attaching the device to the skin, a drug-containing
 layer, and a drug impermeable backing layer. In the
 matrix type system, the drug is dispersed within a solid
 or semi-solid carrier. In the reservoir type system, the
35 drug, typically in liquid solution, is contained within a

walled container, usually formed between the backing layer and a drug-permeable membrane layer.

A number of reservoir type systems have been described. U.S. Patent No. 4,710,191 to Kwiatek et al. involves a reservoir type device that includes a peelable heat sealed release liner layer underlying the reservoir and the portion of the backing layer that extends beyond the reservoir. In a second embodiment, a microporous membrane is included between the backing layer and the reservoir.

U.S. Patent No. 4,829,224 to Chang et al. describes a device with a reservoir that is defined by a backing layer and a drug-permeable membrane layer. A ring-shaped layer made of an adhesive is peripheral to the reservoir. A peelable liner layer underlies the membrane. A second peelable layer, the release liner, underlies the entire assembly. A first heat seal connects the backing layer and the membrane and surrounds the reservoir. A second heat seal concentric about the first heat seal connects the backing layer and the release liner. The second heat seal is broken when the release liner is removed. The device may include an inner liner that underlies the membrane and portions of the backing layer. This inner liner is removed following removal of the release liner so that the membrane is exposed.

U.S. Patent No. 4,983,395 to Chang et al. relates to another device with a backing layer and a membrane layer that define a reservoir. A peelable inner liner underlies the reservoir and portions of the backing and membrane layers outside the periphery of the reservoir. An adhesive layer underlies the inner liner and remaining portions of the backing and membrane layers. A peelable release liner underlies the adhesive layer. A first heat seal connects the backing and

membrane layers on the periphery of the reservoir. A second heat seal underlies the first heat seal and connects the membrane and the inner liner. In use, the release liner and inner liner are peeled away to expose the undersurfaces of the membrane and adhesive layers
5 prior to placement of the device onto the skin or mucosa.

The present invention is an improved device for transdermal or transmucosal drug delivery in which an occlusive peel seal layer underlies the drug containing
10 reservoir and serves to prevent release of the drug prior to use. A release liner overlays the peel seal layer and exposed portions of the adhesive overlay. In this way, the release liner need not be occlusive and thus may be chosen from a wide variety of materials that can be
15 breathable, and can have a stretchable, elastic quality. Such devices are cost efficient and easy to manufacture.

In addition, in the devices of this invention only the central portion of the device in line with the
20 reservoir is occlusive, leaving the portion of the device peripheral to the central portion nonocclusive or breathable.

Disclosure of the Invention

25 The invention is a device for administering an active agent to the skin or mucosa of an individual comprising a laminated composite of:

- (a) an adhesive overlay having a central portion and a peripheral portion;
- 30 (b) a backing layer underlying the central portion of the adhesive overlay;
- (c) an active agent-permeable membrane underlying the backing layer, the backing layer and membrane defining

(d) a reservoir therebetween that contains an active agent;

(e) a peel seal disc underlying the active agent-permeable membrane;

5 (f) a heat seal about the periphery of the peel seal disc, the active agent-permeable membrane and the backing layer; and

(g) a removable release liner underlying the peripheral portion of the adhesive overlay and the peel
10 seal disc.

Brief Description of the Drawings

Figure 1 is an enlarged sectional view of one embodiment of the device of the invention.

15 Figures 2A and 2B are enlarged top views of further embodiments of the device of the invention.

Figure 3 is an enlarged sectional view of another embodiment of the invention.

20 Figure 4 is an enlarged view of the embodiment of Figure 3 showing the embodiment disassembled.

Figure 5 is a graph showing a comparison of calculated and experimental cumulative release kinetics of keterolac tromethamine based on the tests described in Example 23.

25

Modes for Carrying Out the Invention

Figure 1 shows a device, generally designated 10 that is designed to administer a formulation of an active agent to the skin or mucosa. The device 10 is a laminated composite. A drug reservoir 12 is formed
30 between an upper backing layer 14 and an underlying drug-permeable membrane layer 16. A peel seal disc 20 is placed beneath the membrane layer 16. The three layers, the backing layer 14, the membrane layer 16 and the peel
35 seal disc 20 are heat sealed together at their periphery

22, the membrane layer 16 being heat sealed to the backing layer 14 and the peel seal disc 20 being heat sealed to the membrane layer 16. A release liner 24 is placed underneath the peel seal disc 20 and extends
5 beyond the peripheral heat seal 22. An adhesive layer 26 overlies the backing layer 14 and the portions of the release liner 24 that extend beyond the peripheral heat seal 22. An uppermost overlay layer 30 overlies the adhesive layer 26. In this embodiment the release liner
10 is removed together with the peel seal disc prior to placing the device on the skin.

Figures 2A and 2B show further embodiments of the device where a tab 32 extends past the heat seal 22 and allows for easy removal of the peel seal disc 20.
15 The tab 32 may extend a short distance past the heat seal 22 as shown in Figure 2A or may extend to the periphery 34 of the device 10 as shown in Figure 2B.

Figures 3 and 4 depict another embodiment, generally designated 40, of the laminated composite of the invention. The device of Figures 3 and 4 differs from that of Figure 1 by the presence of a second
20 adhesive layer that underlies the peel seal disc and the peripheral portion of the first adhesive layer. More specifically, device 40 consists of (from top surface to basal surface) a nonocclusive overlay layer 42, a first
25 adhesive layer 44, a backing layer 46, a drug reservoir 48 comprising a solution or gelled formulation of a transdermally administrable drug; a drug-permeable membrane layer 50; a peel seal disc 52, a second adhesive
30 layer 54, and a release liner layer 56. As shown, the backing film, reservoir, membrane, and peel seal disc are "in-line" and have smaller areas than the overlay, the two adhesive layers and the release liner. The latter extend peripherally beyond the entire periphery of the
35

former. The backing film, membrane, and peel seal disc are heat sealed at 60 about their entire peripheries.

In use the release liner layer, the portion of the second adhesive layer that underlies the peel seal disc, and the peel seal disc are separated from the rest of the assembly. The separated or disassembled configuration is shown in Figure 4. In this regard, the relative bond strengths between the release layer and the second adhesive layer, the second adhesive layer and the first adhesive layer and peel seal disc, and the peel seal disc and membrane are such that when the release liner is removed, the central portion of the second adhesive layer and the peel seal disc are removed with it. The removed release layer subassembly is discarded. The subassembly containing the drug reservoir is placed on the skin, with the peripheral ring of the second adhesive layer being the means by which that subassembly is affixed to the skin or mucosa. When so placed on the skin the drug reservoir is in diffusional communication with the skin via the membrane. In other words, drug is free to diffuse through the membrane to the skin.

The backing layer 14 or 46 of the device may be composed of a single film or a plurality of films. In any event, its inner surface must be capable of being heat sealed to the membrane layer 16 or 50. One or more of the films that constitute the backing layer will be impermeable to components of the drug formulation contained in the reservoir. Examples of materials used as backing layers in transdermal delivery devices that may find use in the present invention are polyethylene, polypropylene, polyvinylchloride, polyethylene terephthalate, ethylene-vinyl acetate copolymers and combinations thereof. The layer may include one or more metal layers and/or one or more fibrous layers. Preferably the backing layer is occlusive.

The term "active agent" or "drug" as used to describe the principal active ingredient of the device intends a biologically active compound or mixture of compounds that has a therapeutic, prophylactic or other beneficial pharmacological and/or physiological effect on the wearer of the device. Examples of types of drugs that may be used in the inventive device are antiinflammatory drugs, analgesics, antiarthritic drugs, antispasmodics, antidepressants, antipsychotic drugs, tranquilizers, antianxiety drugs, narcotic antagonists, antiparkinsonism agents, cholinergic agonists, anticancer drugs, immunosuppression agents, antiviral agents, antibiotic agents, appetite suppressants, antiemetics, anticholinergics, antihistamines, antimigraine agents, coronary, cerebral or peripheral vasodilators, hormonal agents, contraceptive agents, antithrombotic agents, diuretics, antihypertensive agents, cardiovascular drugs, and the like. The appropriate drugs of such types are capable of permeating through the skin either inherently or by virtue of treatment of the skin with a percutaneous absorption enhancer. Because the size of the device is limited for patient acceptance reasons, the preferred drugs are those that are effective at low concentration in the blood stream. Examples of specific drugs are steroids such as estradiol, progesterone, norgestrel, levonorgestrel, norethindrone, medroxyprogesterone acetate, 3-ketodesogestrel, testosterone and their esters, nitro-compounds such as nitroglycerine and isosorbide nitrates, nicotine, chlorpheniramine, terfenadine, triprolidine, hydrocortisone, oxicam derivatives such as piroxicam, ketoprofen, mucopolysaccharidases such as thiomucase, buprenorphine, fentanyl, naloxone, codeine, dihydroergotamine, pizotiline, salbutamol, terbutaline, prostaglandins such as misoprostol and enprostil, omeprazole, imipramine,

benzamides such as metoclopramine, scopolamine, peptides
such as growth releasing factor and somatostatin,
clonidine, dihydropyridines such as nifedipine,
verapamil, ephedrine, pindolol, metoprolol,
5 spironolactone, nicardipine hydrochloride, calcitriol,
thiazides such as hydrochlorothiazide, flunarizine,
sydononimines such as molsidomine, sulfated
polysaccharides such as heparin fractions and the salts
of such compounds with pharmaceutically acceptable acids
10 or bases.

In addition to the drugs described above,
depending on the permeability of the skin to the drug or
drugs, the reservoir may also contain a percutaneous
absorption enhancer that increases the permeability of
15 the skin to the drug or drugs and is coadministered to
the skin. Examples of percutaneous absorption enhancers
are those referred to in U.S. Patents Nos. 3,989,816,
4,863,970, 4,316,893, 4,405,616, 4,060,084 and 4,379,454
and *J. Pharm Sci* (1975) 64:901-924. The formulation
20 contained in the reservoir may further include solvents,
gelling agents, stabilizers, antiirritants, and other
additives.

The membrane layer 16 or 50 is permeable to the
drug. It may be a "dense" membrane made of material that
25 is inherently permeable to the components of the
reservoir that are to be administered to the skin or
mucosa or it may be made of a microporous membrane whose
pores are filled with a drug-permeable material including
the drug reservoir formulation itself which may include
30 enhancers where desired. In the case of dense membranes,
the components dissolve in the material and diffuse
through the material to the skin. In the case of
microporous materials, the components diffuse through the
pores to the skin. The membrane may or may not be a
35 rate-controlling element depending upon the particular

drug involved, the permeability of the skin to the drug, and the rate of delivery required to provide therapy. Examples of materials for making dense membranes are given in U.S. Patents Nos. 3,598,122 and 4,650,484.

- 5 Examples of materials for making microporous membranes are provided in U.S. Patents Nos. 3,797,494 and 4,031,894.

The adhesive layers 26, 44 and 54 are composed of pressure sensitive surgical adhesive such as those
10 that are commonly used to affix transdermal drug delivery devices, bandages or other dressings to the skin. Examples of such adhesives are polyisobutene (PIB), natural rubber adhesives, acrylate and methacrylate adhesives and silicone adhesives. Breathable,
15 nonocclusive adhesives such as the acrylate/methacrylate and silicone adhesives are preferred.

The peel seal disc 20 or 52 may be composed of a single layer or a multiplicity of layers. The disc should be (1) impermeable to the components of the drug
20 reservoir formulation that diffuse through the membrane, (2) heat-sealable to the membrane layer, and (3) inherently strippable or rendered so by techniques such as silicone or fluorocarbon treatment or surface treatment with a seal incompatible layer. Suitable peel
25 seal discs include those made from Bertek 4418 Peelable Seal, Total Healthcare Packaging TPC-0812 or TPC-0760 and UCB Medical Industries LR 4/25.

The release liner 24 or 56 may similarly be composed of a single layer or a multiplicity of layers.
30 Unlike the peel seal disc, however, the release liner need not be impermeable to the components of the drug formulation in that the peel seal disc protects against release of the components from the reservoir. Therefore, suitable release liners may be made from materials such
35 as polyester, low density polyethylene (LDPE), high

density polyethylene (HDPE), polypropylene, polystyrene, polyamide, nylon, polyvinyl chloride and specialty papers, and include Akrosil Biorelease liners, Scotchpak 1022 release liners, Adhesives Research AR5MS, Custom
5 Coating and Laminating 7000 on HDPE or 6020 on polyethylene terephthalase (PET).

The overlay layer 30 or 42 overlies the adhesive layer 26 or 44 and may be made from materials such as polyolefin, polyurethane, nylon, polyester,
10 vinyl, acetate taffeta or other elastomeric woven or nonwoven fabrics or films. Preferably the overlay material is nonocclusive (breathable).

The respective components of the device may be formulated and assembled using procedures that are known
15 in the drug formulation, transdermal device and laminating arts. The shape of the device is not critical, and devices of preformed shapes may be assembled directly or punched, cut or otherwise formed from large sheets of laminated composite.

20 The following examples further illustrate the invention. These examples are not intended to limit the invention in any manner.

EXAMPLES

25

Example 1

A medical grade, pressure sensitive acrylic adhesive, MA-31 (Adhesives Research, Glen Rock, PA) was coated onto a low peel strength, siliconized release
30 liner using a 10 mil gap casting knife. The adhesive solvent was evaporated at 80°C for 15 min. to provide a final dry adhesive thickness of .002 inches. Volara Foam 15EO (Voltek, Division of Sekisui America Corp, Lawrence MA) the overlay material, was then laminated onto the dry
35 adhesive film forming the overlay adhesive laminate.

This overlay adhesive laminate was used at a later stage of the fabrication process as described below.

A gelled testosterone formulation was prepared by dissolving 15 mg/ml testosterone in a solution
5 containing 50.0/15.0/30.0/2.5/2.5% (v/v) of ethanol/water/glycerin/glycerol monooleate/methyl laureate. The resulting solution was gelled with 3.0% (w/v) of Carbopol 1342. The pH of the resulting gel was adjusted to 4.5 to 5.0 by addition of 2N sodium
10 hydroxide.

The overlay patch with peel seal disc was fabricated as follows: A CoTran 9711 microporous membrane (3M, St. Paul, MN) was placed over an LR4/25 peel seal disc film (UCB Medical Industries, Bloomfield,
15 CT). 0.9 g of the testosterone formulation was dispensed onto the microporous membrane. The backing film (Scotchpak 1012, 3M, St. Paul MN) was predimpled to provide a flat, circular cup with a volume of approximately 1.0 cc. This predimpled backing film was
20 placed over the gel such that the dimple in the backing film was centered on the mound of gel. The backing film was then heat sealed to the microporous membrane/peel seal film laminate at 370°F with a pressure of 30 psi and a 0.4 sec. dwell time. A circular heat seal die with a
25 interior diameter of 1.22 inches and a heat seal width of 0.03 inch was used. A gel containing reservoir with an active surface of 7.5cm² was formed. This heat sealing step simultaneously creates a permanent reservoir seal between the microporous membrane and the backing film and
30 a peelable seal between the microporous membrane and the peel seal disc film. This intermediate drug containing reservoir system, with a diameter of 3.5 cm and a total surface area of 9.6 cm², was then diecut from the heat sealed laminate.

The above reservoir system was laminated to the previously prepared overlay adhesive laminate by removing the protective release liner from the overlay adhesive, placing the diecut reservoir system onto the adhesive with the backing film contacting the adhesive, relaminating the release liner over the exposed adhesive and finally diecutting the actual overlay adhesive transdermal delivery system from this laminate. The cutting die used for this operation has a diameter of 5.5 cm thereby diecutting a transdermal delivery system with a final surface area of 23.8 cm², which includes the peripheral adhesive area. The activation of this system before application to the skin proceeds in a two step process: the release liner covering the adhesive is removed first, followed by the removal of the peelable disc film protecting the reservoir.

Example 2

Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the adhesive used was a medical grade acrylic adhesive, Gelva 737 (Monsanto, St. Louis, MO).

Example 3

Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the adhesive used was a medical grade PIB adhesive MA-24 (Adhesives Research, Glen Rock, PA).

Example 4

Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that a silicone based adhesive, BIO PSA X7-2920 (Dow Corning, Midland, MI) was used. The release liner

used was the Akrosil BioRelease liner (Menasha, WI) which is compatible with the above silicone adhesive.

Example 5

5 Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the adhesive used was an (acrylic copolymer) adhesive, TSR Adhesive (Sekisui Chemical Company, Osaka, Japan).

10

Example 6

Transdermal systems with an adhesive overlay were made using the same procedure as described in Example 1, except that the peel seal disc film was a
15 multilaminate PET/LDPE/Foil/Primacor 3440/LLDPE/Heat Seal Film flexible packaging material, TPC-0812 (Tolas Healthcare Packaging, Feasterville, PA).

Example 7

20 Transdermal systems with an adhesive overlay were made using the same procedure as described in Example 1, except that the peel seal disc film was a Peelable Foil Film material, a multilaminate of PET/adhesive/Foil/adhesive/LLDPE/Heat Seal Film
25 (Technipaq, Crystal Lake, IL).

Example 8

Transdermal systems with an adhesive overlay were made using the procedure described in Example 1,
30 except that the overlay material was an embossed polyurethane elastomeric film.

Example 9

Transdermal systems with an adhesive overlay
35 were made using the procedure described in Example 1,

except that the overlay material was a 100% nonwoven polyester fabric.

Example 10

5 Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the overlay material was a spunbonded nylon film.

Example 11

10 Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the overlay material was a woven acetate taffeta film.

Example 12

15 A 4 in. x 4 in. commercially available wound dressing, Mitraflex[®] Plus, distributed by Calgon Vestal Laboratories, was used as the adhesive overlay.

20 A gelled ketorolac tromethamine formulation was prepared by dissolving 60 mg/ml ketorolac tromethamine in a solution containing 55.28/33.17/10.0/1.55% (v/v) of isopropyl alcohol/water/glycerin/isopropyl myristate. The resulting solution was gelled with 5.0% (w/v) of
25 Natrosol[®] Plus 330 CS and the pH of the resulting gel was adjusted to $5.1 \pm (0.2)$ by addition of 2N HCl. The gel also contained 0.1% (w/v) butylated hydroxy toluene as an antioxidant.

The overlay patch with peel seal disc was
30 fabricated as follows: A CoTran 9710 microporous membrane was placed over an LR4/25 peel seal disc film. 3.44 g of the ketorolac tromethamine formulation was dispensed onto the microporous membrane. The backing film was predimpled to provide a flat, circular cup.
35 This predimpled backing film was placed over the gel such

that the dimple in the backing film was centered on the mound of gel. The backing film was then heat sealed to the microporous membrane/peel seal film laminate at 370°F with a pressure of 30 psi and a 0.4 sec. dwell time. An
5 oval heat seal die was used to form a gel containing reservoir with an active surface of 30 cm². This heat sealing step simultaneously creates a permanent reservoir seal between the microporous membrane and the backing film and a peelable seal between the microporous membrane
10 and the peel seal disc film. This intermediate drug containing reservoir system, with a total surface area of 42.55 cm², was then diecut from the heat sealed laminate.

The above reservoir systems were laminated to the Mitraflex[®] Plus overlay adhesive laminate by removing
15 the protective release liner from the wound dressing, placing the diecut reservoir system onto the adhesive with the backing film contacting the adhesive and relaminating the release liner over the exposed adhesive to obtain the actual overlay adhesive transdermal
20 delivery system. The activation of this system before application to the skin proceeds in a two step process: the release liner covering the adhesive is removed first, followed by the removal of the peelable disc film protecting the reservoir.

25

Example 13

Transdermal systems with an adhesive overlay were made using the procedure described in Example 12, except that the overlay adhesive laminate used was
30 fabricated as in Example 1. The final surface area of the transdermal delivery system was 63 cm².

Example 14

Transdermal systems with an adhesive overlay
35 were made using the procedure described in Example 13,

except that the adhesive used was a medical grade PIB adhesive, MA-24 (Adhesives Research, Glen Rock, PA).

Example 15

5 Transdermal systems with an adhesive overlay were made using the procedure described in Example 13, except that a silicone based adhesive, BIO PSA X7-2920 (Dow Corning, Midland, MI). The release liner used was the Akrosil BioRelease liner (Menasha, WI) which is
10 compatible with the above silicone adhesive.

Example 16

Transdermal systems with an adhesive overlay were made using the procedure described in Example 13,
15 except that the adhesive used was an acrylic copolymer adhesive, TSR Adhesive (Sekisui Chemical Company, Osaka, Japan).

Example 17

20 Transdermal systems with an adhesive overlay were made using the procedure described in Example 13, except that the peel seal disc film was a multilaminate of PET/LDPE/Foil/Primacor 3440/LLDPE/Heat Seal film flexible packaging material, TPC-0812 (Tolas Healthcare
25 Packaging, Feasterville, PA).

Example 18

Transdermal systems with an adhesive overlay were made using the same procedure as described in
30 Example 13, except that the peel seal disc film was a Peelable Foil Film, a multilaminate PET/adhesive/Foil/adhesive/LLDPE/Heat Seal Film (Technipaq, Crystal Lake, IL).

Example 19

Transdermal systems with an adhesive overlay were made using the procedure described in Example 13, except that the overlay material was an embossed polyurethane elastomeric film.

Example 20

Transdermal systems with an adhesive overlay were made using the procedure described in Example 13, except that the overlay material was a 100% nonwoven polyester fabric.

Example 21

Transdermal systems with an adhesive overlay were intermediate drug-c made using the procedure described in Example 13, except that the overlay material was a spunbonded nylon film.

Example 22

Transdermal systems with an adhesive overlay were made using the procedure described in Example 13, except that the overlay material was a woven acetate taffeta film.

Example 23

Ketorolac tromethamine gel composition was prepared as described in Example 12. The gel was evaluated for in vitro skin flux performance.

In vitro human cadaver skin flux studies were conducted using modified Franz diffusion cells, placed in a water bath calibrated to maintain the skin surface temperature at 32°C. The epidermal membrane was separated from the full thickness human cadaver skin by the method of Kligman and Christopher (Arch. Dermatol. 88, 702-705, 1963). The full thickness skin was exposed

to 60°C heat for 60 seconds. The stratum corneum and epidermis were gently peeled from the dermis. The epidermal membrane was mounted onto a 0.65 cm² surface area diffusion cell with the epidermal side facing the receiver compartment and clamped in place. The receiver compartment was then filled with an aqueous solution containing 0.02% sodium azide as a bacteriostat. The cell was placed in a circulating water bath calibrated to maintain skin surface temperature at 32±1°C and the skin was allowed to hydrate overnight. The subsequent morning, 75µl of the gelled ketorolac formulation was pipetted into a cavity created by placing a Teflon washer over the stratum corneum surface. The cavity was then occluded by clamping an occlusive backing film over the Teflon washer assembly. Throughout the experiment, the receiver compartment, which contained a bar magnet, was continuously stirred by a magnetic stirrer placed under the water bath. At predetermined intervals, the entire contents of the receiver compartment was collected for drug quantitation using the HPLC method in Table I. The receiver compartment was refilled with fresh receptor medium, taking care to eliminate any air bubbles at the skin/solution interface. From the concentration versus time curve, the average steady state flux, the interval flux and cumulative delivery per unit area at 24 hours were calculated. The in vivo delivery was projected by multiplying the in vitro cumulative delivery (µg/cm²/24h) by the patch active area (30 cm²).

30

35

Table I
In Vitro Skin Flux
HPLC Method

5	Column:	Partisphere C-18 (100 mm)
	Flow Rate:	1.0 ml/min
	Wavelength:	314 nm
10	Injection Volume:	10 μ l
	Retention Time:	2.0 \pm 0.3 min
15	Mobile Phase:	60/40/0.5 (%v/v) Acetonitrile/H ₂ O/Glacial Acetic Acid pH ~ 3.0

Overlay patches, prepared as described in
Example 12, were applied to the chest in 24 healthy adult
volunteers (12 male and 12 female) in human clinical
trials. Plasma samples were collected for ketorolac
quantitation at predetermined times. After removal of
the patches in each study, the used patches were
extracted in 100 ml of methanol and analyzed for drug
content using the HPLC method described in Table II. The
difference between the initial and final patch drug
contents gave the amount of ketorolac delivered, based on
patch depletion analysis.

30

35

Table II
Patch
Content Assay
HPLC Method

5	Column:	Zorbax RX-C8 5 μ m, 15 cm x 4.6 mm
	Flow Rate:	1.0 ml/min
10	Wavelength:	314 nm
	Injection Volume:	10 μ l
	Retention Time:	5.2 \pm 0.2 min
15	Mobile Phase:	60/40/1 (%v/v) Acetonitrile/H ₂ O/Glacial Acetic Acid pH ~ 3.0

Table III compares the in vivo cumulative delivery, based on patch depletion analysis, with the projected drug delivery for the 30 cm² overlay patch, based on in vitro skin flux results. The projected drug delivery is based upon average in vitro cumulative deliver (μ g/cm²/24h) multiplied by the active area (30 cm²) of the overlay patch. The average in vivo dose was calculated from the difference between initial drug content of the patches and the residual drug content of the patches upon removal. In vivo, the 30 cm² overlay patch delivered (46.4 \pm 17.5) mg/day of ketorolac tromethamine, in very good agreement with the predicted value of 53.7 mg/day. Figure 5 shows that the average in vivo cumulative input kinetics, calculated according to the method of Wagner and Nelson (J. Pham. Sci. 52, 610-611, 1963) is in very good agreement with the in vitro cumulative input kinetics.

Table III

Projected and Observed
In Vivo Performance
of Overlay Patch System

5

10	Patch System	Projected Dose Ketorolac Tromethamine (mg/day)	In Vivo Dose Ketorolac Tromethamine (mg/day)
	30 cm ² Overlay Patch	53.7	46.4 ± 17.5 (24 subjects)

15

Example 24

Transdermal systems with an adhesive overlay were made using the procedure described in Example 1, except that the die used to diecut the liquid reservoir portion of the transdermal delivery system allowed the peel seal disc tab to reach the edge of the release liner, as shown in Figure 2B. This extended tab allows the final system to be activated in a single step process, instead of the two step process described in Example 1. In the single step process, the release liner and the extended tab of the peel seal disc are held between the thumb and index finger of one hand and the backing film in the other. The peel seal disc and release liner are pulled apart from the backing film to activate the system.

30

Example 25

A medical grade pressure sensitive adhesive, MA-31 (Adhesives Research, Glen Rock, PA) was coated onto a high peel strength, siliconized release liner using a 10 mil gap casting knife. The adhesive solvent was

35

evaporated at 80°C for 15 min. to provide a final dry adhesive thickness of .0015 inches. A low strength, in-process, 2 mil high density polyethylene silconized liner was then laminated onto the dry adhesive film forming the primary adhesive laminate. This primary adhesive laminate, and the overlay adhesive laminate fabricated as described in Example 1, were used at a later stage of the fabrication process as described below.

The intermediate drug containing reservoir was fabricated as an Example 1, except that the weight of the testosterone gel was 1.8g, the active surface was 15 cm², and the total surface area was 18.4 cm². This reservoir system was laminated to the primary adhesive laminate by removing the in-process liner, placing the diecut reservoir centrally onto the adhesive with the peel-seal disc film contacting the adhesive. The protective liner from the overlay adhesive laminate was then removed and the overlay adhesive laminated onto the primary adhesive/reservoir system, such that the overlay adhesive was in contact with the backing film of the intermediate drug-containing reservoir and also the adhesive area of the primary adhesive laminate extending beyond the intermediate drug-containing reservoir. The final overlay adhesive transdermal delivery system was diecut from this laminate structure to give a system with a total surface area of 33 cm².

The activation of this system is carried out in a single step by pulling away the high peel strength liner from the overlay adhesive. The peel-seal disc remains anchored to the liner, thereby exposing the reservoir area.

Example 26

Transdermal systems with an adhesive overlay were made using the same procedure as described in Example 25, except that the adhesive used in the primary adhesive laminate and the overlay adhesive laminate was a medical grade polyisobutene (PIB) adhesive MA-24 (Adhesives Research, Glen Rock, PA).

10

Example 27

Transdermal systems with an adhesive overlay were made using the same procedure as described in Example 25, except that the adhesive used in the primary adhesive laminate and the overlay adhesive laminate was a silicone based adhesive, BIO PSA X7-2920 (Dow Corning, Midland, MI).

15

Modifications of the above-described modes for carrying out the invention that are obvious to those of skill in the fields of chemistry, transdermal drug delivery, pharmacology and related fields are intended to be within the scope of the following claims.

20

25

30

35

CLAIMS

We claim:

1. A device for administering an active agent
5 to the skin or mucosa of an individual comprising a laminated composite of:
 - (a) an adhesive overlay having a central portion and a peripheral portion;
 - (b) a backing layer underlying the central
10 portion of the adhesive overlay;
 - (c) an active agent-permeable membrane underlying the backing layer, the backing layer and membrane defining
 - (d) a reservoir therebetween that contains a
15 formulation of the active agent;
 - (e) a peel seal disc underlying the active agent-permeable membrane;
 - (f) a heat seal about the periphery of the peel seal disc, the active agent-permeable membrane and
20 the backing layer; and
 - (g) a removable release liner underlying the peripheral portion of the adhesive overlay and the peel seal disc.
- 25 2. The device of claim 1 wherein the adhesive is made of a material selected from the group consisting of polyisobutenes, natural rubbers, acrylates, methacrylates and silicone.
- 30 3. The device of claim 1 wherein the backing layer comprises at least one impermeable film layer.
4. The device of claim 1 wherein the membrane
35 is selected from the group consisting of microporous membranes and dense membranes.

5. The device of claim 1 wherein the peel seal disc is composed of at least one layer.

6. The device of claim 1 further comprising
5 a percutaneous absorption enhancer.

7. The device of claim 1 whenever the formulation further comprises a material selected from the group consisting of a solvent, a gelling agent,
10 stabilizer and an anti-irritant.

8. The device of claim 1 including an adhesive layer that underlies the peripheral portion of the adhesive overlay and the peel seal disc and overlies the
15 release liner.

9. The device of claim 1 wherein the adhesive overlay is nonocclusive and the backing layer is occlusive.
20

10. A device for administering an active agent to the skin or mucosa of an individual comprising a laminated composite of:

- (a) an adhesive overlay having a central
25 portion and a peripheral portion;
- (b) a backing layer underlying the central portion of the adhesive overlay;
- (c) an active agent-permeable membrane, the backing layer and membrane defining
30
- (d) a reservoir therebetween that contains a formulation of the active agent;
- (e) a peel seal disc underlying the active agent-permeable membrane;

(f) a heat seal about the periphery of the peel seal disc, the active agent-permeable membrane and the backing layer; and

(g) a removable release liner underlying the peripheral portion of the adhesive overlay and the peel seal disc; and

(h) a tab on said peel seal disc to aid in removal of said disc.

10 11. The device of claim 10 including an adhesive layer that underlies the peripheral portion of the adhesive overlay and the peel seal disc and overlies the release liner.

15 12. The device of claim 10 wherein the adhesive overlay is nonocclusive and the backing layer is occlusive.

20

25

30

35

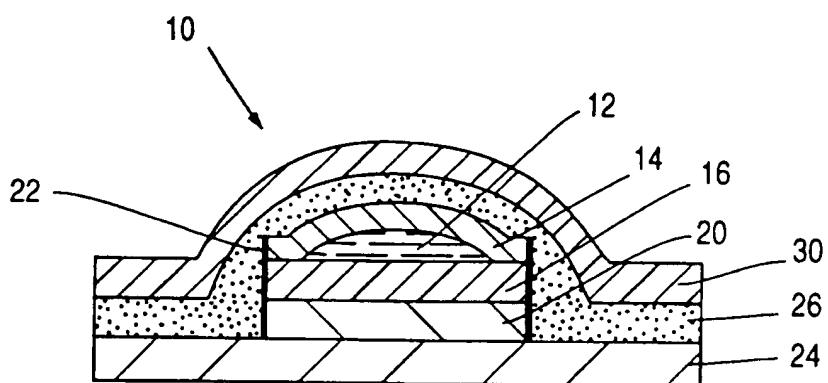
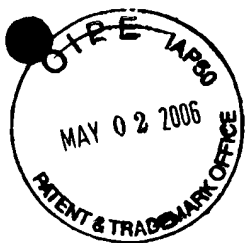


FIG. 1

2/4

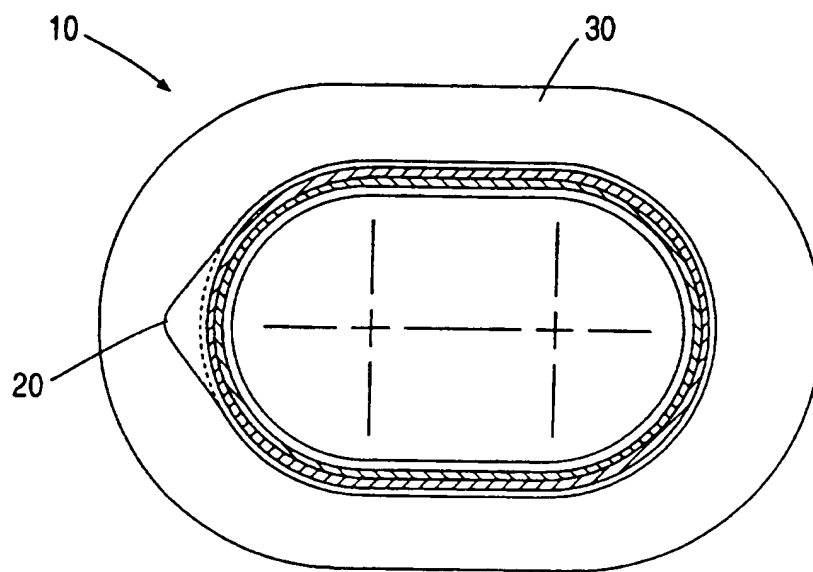


FIG. 2A

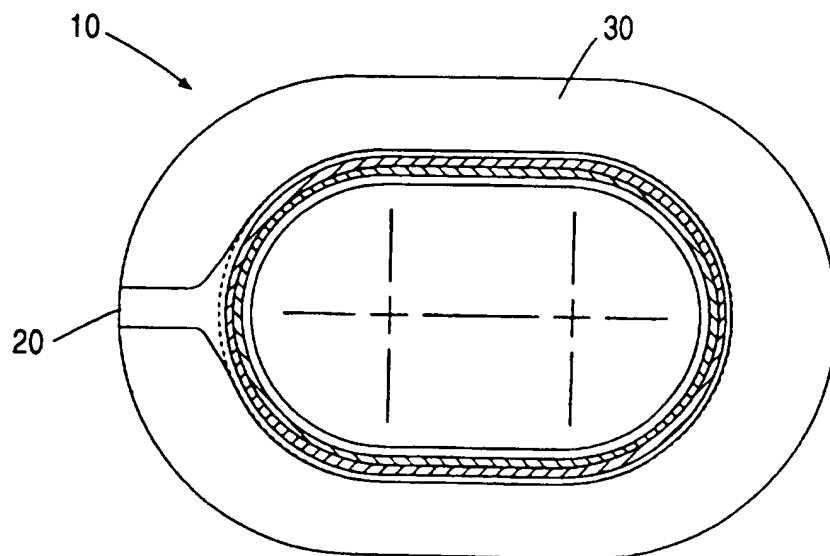


FIG. 2B

3/4

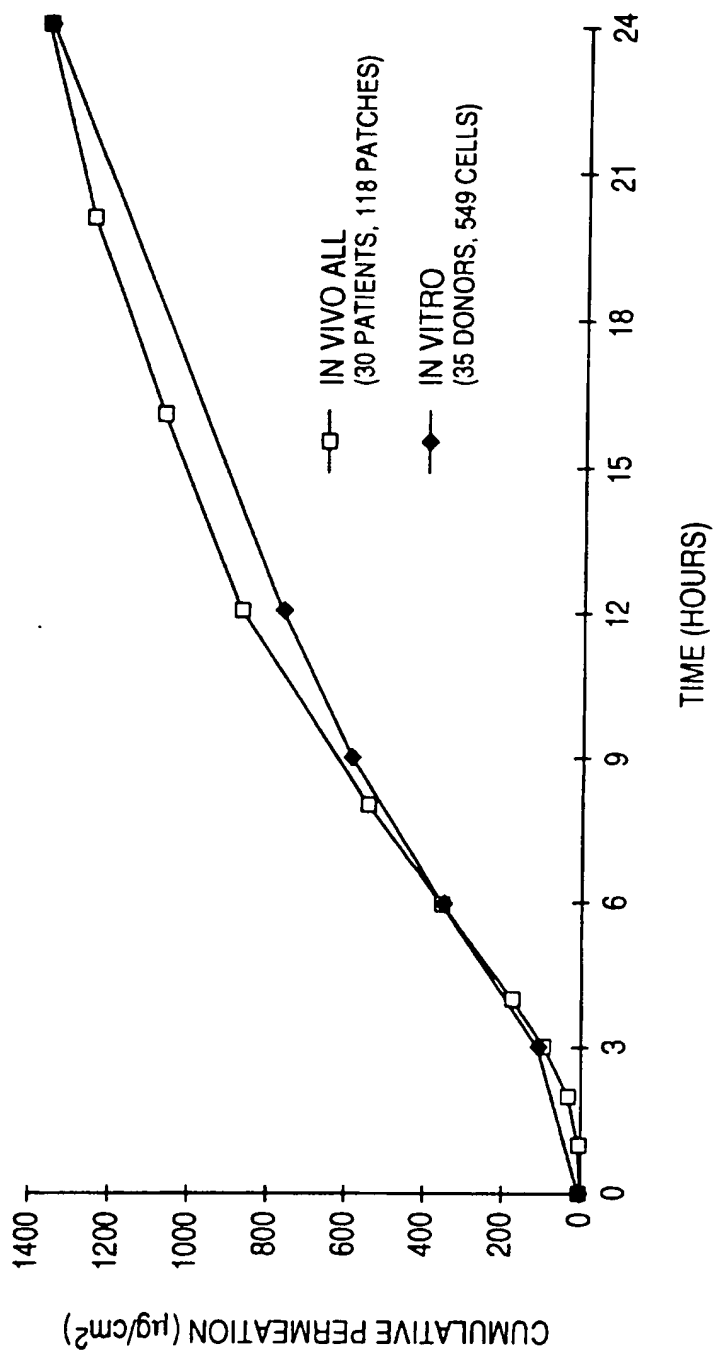


FIG. 3

4/4

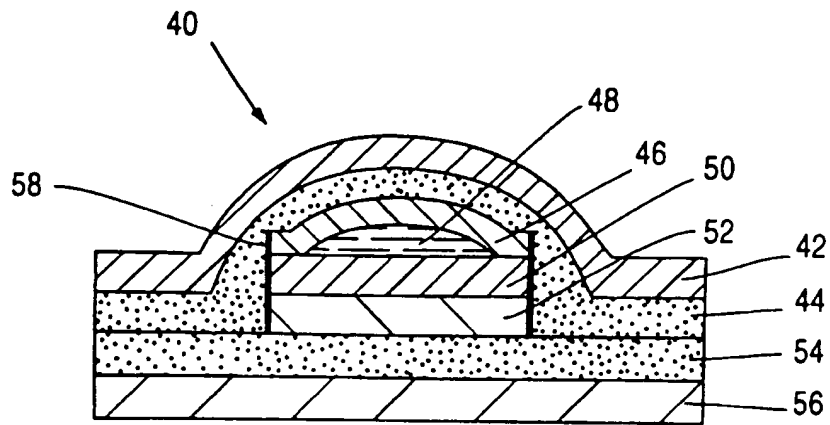


FIG. 4

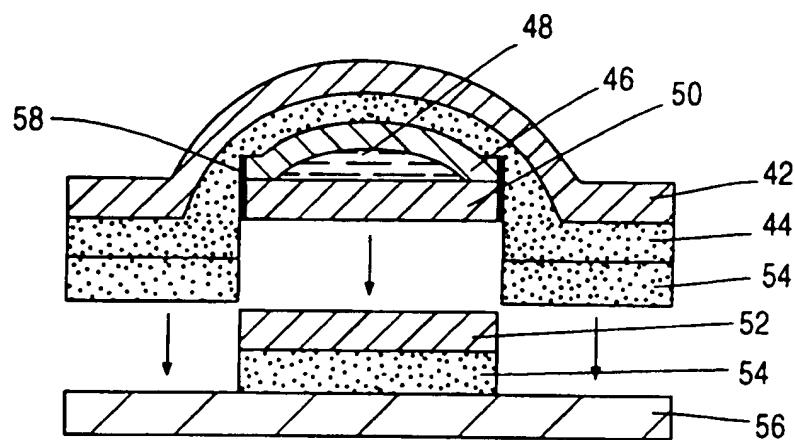


FIG. 5

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 95/16498

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61K9/70

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO,A,92 10154 (THERATECH, INC.) 25 June 1992 see page 20, line 5 - line 22 see figure 2 ---	1-12
Y	GB,A,2 184 016 (JONERGIN INC.) 17 June 1987 see page 1, line 121 - page 2, line 84 see figure 1 & US,A,4 710 191 cited in the application ---	1-12
A	EP,A,0 040 861 (KEY PHARMACEUTICALS) 2 December 1981 see page 12, line 33 - page 14, line 2 see figure 2 --- -/-	1-12

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *&* document member of the same patent family

Date of the actual completion of the international search

8 May 1996

Date of mailing of the international search report

15.05.96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl,
Fax (+ 31-70) 340-3016

Authorized officer

Benz, K

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/US 95/16498

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>WO,A,93 23019 (SRI INTERNATIONAL) 25 November 1993 see page 9, line 20 - line 27 see figure 1 -----</p>	1-12

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 95/16498

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO-A-9210154	25-06-92	US-A- 5202125	13-04-93
		US-A- 5122383	16-06-92
		AU-B- 656755	16-02-95
		AU-B- 9141391	08-07-92
		CA-A- 2098196	10-06-92
		EP-A- 0561983	29-09-93
		JP-T- 6503576	21-04-94
		US-A- 5302395	12-04-94
		US-A- 5227169	13-07-93
US-A- 5212199	18-05-93		

GB-A-2184016	17-06-87	US-A- 4710191	01-12-87
		BE-A- 905928	16-04-87
		CA-A- 1261702	26-09-89
		CH-A- 670393	15-06-89
		DE-A- 3642600	19-06-87
		FR-A- 2591493	19-06-87
		JP-A- 62195326	28-08-87
		NL-A- 8603204	16-07-87
		SE-C- 500637	01-08-94
		SE-A- 8605320	17-06-87

EP-A-40861	02-12-81	AT-T- 7266	15-05-84
		AU-B- 5445280	17-07-80
		CA-A- 1135625	16-11-82
		EP-A, B 0013606	23-07-80
		EP-A- 0040862	02-12-81
		JP-B- 1021128	19-04-89
		JP-C- 1539119	16-01-90
		JP-A- 55094316	17-07-80
		US-A- 4291014	22-09-81
		US-A- 4579731	01-04-86

WO-A-9323019	25-11-93	NONE	
